

Used in Lieu of PTO/SB/08A/B
(Based on PTO 04-07 version)

Substitute for form 1449 INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)				Complete If Known	
				Application Number	10/767,018
				Filing Date	January 29, 2004
				First Named Inventor	Brent R. Stockwell
				Art Unit	1643
				Examiner Name	K. A. Canella
Sheet	1	of	1	Attorney Docket Number	WIBL-P01-011

U.S. PATENT DOCUMENTS					
Examiner Initials*	Cite No. ¹	Document Number	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number-Kind Code ² (if known)			
/KAC/	AA	US-6,831,085	12-14-2004	Bergnes et al.	

FOREIGN PATENT DOCUMENTS						
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		Country Code ³ -Number ⁴ -Kind Code ⁵ (if known)				
/KAC/	BA	JP-07-258224-A	10-09-1995	Dai Ichi Seiyaku Co. Ltd (abstract)		

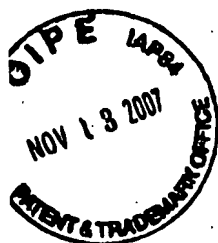
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/KAC/	AB	US-20040248221-A1	12-09-2004	Stockwell	
/KAC/	AC	US-20030171316-A1	09-11-2003	Jupe	
/KAC/	AD	US-20040096444-A1	05-20-2004	Pizzo et al.	

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		Country Code ³ -Number ⁴ -Kind Code ⁵ (if known)				
/KAC/	BB	WO-02/099122	12-12-2002	Exelixis Inc et al.		
/KAC/	BC	WO-2004/030615	04-15-2004	Genentech Inc et al.		
/KAC/	BD	WO-04/055519	07-01-2004	Hoffmann La Roche et al.		
/KAC/	BE	WO-99/21988	05-06-1999	Shanghai Second Medical Univer et al.		
/KAC/	BF	WO-02/083143	10-24-2002	Tularik Inc et al.		

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/KAC/	CR2	ABDEL-ALIM, et al., "Synthesis and biological activities of 6-bromo-2,3-disubstituted-4-(3H)-quinazolinones," Indian Journal of Chemistry, 33(B):260-265 (1994).				
/KAC/	CS2	ADAM, et al., "Comprehensive Proteomic Analysis of Breast Cancer Cell Membranes Reveals Unique Proteins with Potential Roles in Clinical Cancer," JBC Papers in Press, 1-60 (2002).				
/KAC/	CT2	AGER, et al., "Synthesis and Central Nervous System Activity of Quinazolones Related to 2-Methyl-3-(o-tolyl)-4(3H)-quinazolinone (Methaqualone)," J. Med. Chem., 20(3):379-386 (1977).				
/KAC/	CU2	Database Registry Chemical Abstracts Service, Columbus, Ohio, US; (2001-05-21), XP002405284, RN 336853-04-4.				
/KAC/	CV2	Database Registry Chemical Abstracts Service, Columbus, Ohio, US; (2001-05-21), XP002405285, RN 336813-90-2.				
/KAC/	CW2	DOLMA, et al., "Identification of genotype-selective antitumor agents using synthetic lethal chemical screening in engineered human tumor cells," Cancer Cell, 3:285-296 (2003).				
/KAC/	CX2	FIGYS, et al., "VDAC Can Control Apoptosis By Controlling Metabolism," Biophysical Jr., 86(1):463A-464A (2004).				
/KAC/	CY2	GUPTA, et al., "A Novel Class of Hypoglycaemic Agents: Syntheses & SAR in 2-Substituted 4(3H)-Quinazolones, 2-Substituted 4-Hydroxypolymethylene[5,6]pyrimidines & 3-Substituted 4-Oxo-pyrido[1,2- α]pyrimidines," Indian Journal of Chemistry, 9:201-206 (1971).				
/KAC/	CZ2	IKONEN, et al., "Prohibitin, an antiproliferative protein, is localized to mitochondria," FEBS Letters, 358(3):273-277 (1995).				
/KAC/	CA3	KOZHEVNIKOV, et al., "Synthesis in the 2-aminoethyl-3-(2'-tolyl)-4-quinazolinone," Khimiko-				
Examiner Signature				Date Considered		

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				First Named Inventor	Brent R. Stockwell
				Art Unit	1643
				Examiner Name	K. A. Canella
Sheet	2	of	2	Attorney Docket Number	WIBL-P01-011

		Farmatsevticheskii Zhurnal, 4(11):22-25 (1970).	
/KAC/	CB3	TANI, et al., "Studies on Biologically Active Halogenated Compounds II. Chemical Modifications of 6-amino-2-fluoromethyl-3-(o-toly)-4(3H) quinazolinone and the CNS depressant activities of related compounds," Chemical and Pharmaceutical Bulletin, Pharmaceutical Society of Japan, 27(11):2675-2687 (1979).	
/KAC/	CC3	VERMA, et al., "A New Potent Anti-Inflammatory Quinazolone," Pharmacological Research Communications, Italian Pharmacological Society, IT, 13(10):967-979 (1981).	

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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)		Application Number	10/767018
		Filing Date	January 29, 2004
		First Named Inventor	Brent R. Stockwell
		Art Unit	1642
		Examiner Name	Not Yet Assigned
		Attorney Docket Number	WIBL-P01-011
Sheet	2	of	4

KAC		chemical screening in engineered human tumor cells," <i>Cancer Cell</i> , 3:285-296 (2003).	
KAC	CN	Druker, B.J. et al., "Effects of a selective inhibitor of the Abl tyrosine kinase on the growth of Bcr-Abl positive cells", <i>Nature Medicine</i> , 2:561-566 (1996) (Abstract)	
	CO	Elenbaas, B. et al., "Human breast cancer cells generated by oncogenic transformation of primary mammary epithelial cells", <i>Genes & Development</i> , 15:50-65 (2001)	
	CP	Eng, W.-K., et al., "Evidence that DNA Topoisomerase I is Necessary for the Cytotoxic Effects of Camptothecin," <i>Mol Pharmacol</i> , 34:755-60 (1988).	
	CQ	Hahn, W. C. and Weinberg, R. A., "Modelling the Molecular Circuitry of Cancer", <i>Nature Reviews Cancer</i> , 2:331-341 (2002)	
	CR	Hahn, W.C., et al., "Creation of human tumour cells with defined genetic elements," <i>Nature</i> , 400:464-468 (1999).	
	CS	Hahn, W.C., et al., "Enumeration of the Simian Virus 40 Early Region Elements Necessary for Human Cell Transformation," <i>Mol Cell Biol</i> , 22(7):2111-23 (2002).	
	CT	Hahn, W.C., et al., "Inhibition of telomerase limits the growth of human cancer cells," <i>Nat Med</i> , 5(10):1164-1170 (1999).	
	CU	Hamad, N. M. et al., "Distinct requirements for Ras oncogenesis in human versus mouse cells", <i>Genes & Development</i> , 16:2045-2057 (2002)	
	CV	Harley, C.B., "Telomerases," <i>Pathol Biol (Paris)</i> , 42:342-5 (1994).	
	CW	Hsiang, Y.-H. and Liu, L.F., "Identification of Mammalian DNA Topoisomerase I as an Intracellular Target of the Anticancer Drug Camptothecin," <i>Cancer Res</i> , 48:1722-6 (1988).	
	CX	Hsiang, Y.-H., et al., "Arrest of Replication Forks by Drug-stabilized Topoisomerase I-DNA Cleavable Complexes as a Mechanism of Cell Killing by Camptothecin," <i>Cancer Res</i> , 49:5077-82 (1989).	
	CY	Jorcyk, C.L., et al., "Development and Characterization of a Mouse Prostate Adenocarcinoma Cell Line: Ductal Formation Determined by Extracellular Matrix," <i>The Prostate</i> , 34:10-22 (1998).	
	CZ	Kohno, T., et al., "Alterations of the PPP1R3 Gene in Human Cancer," <i>Cancer Res</i> , 59:4170-4 (1999).	
	CA1	Laurent, G. and Jaffrezou, J.-P., "Signaling pathways activated by daunorubicin," <i>Blood</i> , 98(4):913-924 (2001).	
	CB1	Lessnick, S.L., et al., "The Ewing's sarcoma oncoprotein EWS/FLI induces a p53- dependent growth arrest in primary human fibroblasts," <i>Cancer Cell</i> , 1:393-401 (2002).	
	CC1	Liu, L.F., et al., "Mechanism of Action of Camptothecin," <i>Annals N Y Acad Sci</i> , 922:1-10 (2000).	
	CD1	Loomis, C.R. and Bell, R.M., "Sangivamycin, a Nucleoside Analogue, Is a Potent Inhibitor of Protein Kinase C*," <i>J Biol Chem</i> , 263(4):1682-1692 (1998).	
	CE1	Madden, K.R., and Champoux, J.J., "Overexpression of Human Topoisomerase I in Baby Hamster Kidney Cells: Hypersensitivity of Clonal Isolates to Camptothecin," <i>Cancer Res</i> , 52:525-32 (1992).	
	CF1	Majno, G. and Joris, I., "Apoptosis, Oncosis, and Necrosis," <i>Am J Pathol</i> , 146(1):3-15 (1995).	
	CG1	Makin, G., "Targeting apoptosis in cancer chemotherapy," <i>Expert Opin Ther Targets</i> , 6(1):73-84 (2002).	
	CH1	Miller, M.L. and Ojima, I., "Chemistry and Chemical Biology of Taxane Anticancer Agents," <i>Chem. Record</i> , 1:195-211 (2001).	
	CI1	Millward, T.A., et al., "Regulation of protein kinase cascades by protein phosphatase 2A," <i>Trends Biochem Sci</i> , 24:186-91 (1999).	
	CJ1	Mokbel, K. and Hassanally, D., "From HER2 to Herceptin," <i>Curr Med Res Opin</i> , 17(1):51-9 (2001).	
↓	CK1	Möller, I., et al., "Anthracycline-derived chemotherapeutics in apoptosis and free radical	

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Under the Paperwork Reduction Act of 1995, no persons are required to respond to a collection of information unless it contains a valid OMB control number.

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KAC		cytotoxicity (Review), <i>Int J Mol Med</i> , 1:491-4 (1998).	
KAC	CL1	Nociari, M.M., <i>et al.</i> , "A novel one-step, highly sensitive fluorometric assay to evaluate cell-mediated cytotoxicity," <i>J. Immunol. Methods</i> , 213:157-167 (1998).	
	CM1	Pallas, D.C., <i>et al.</i> , "Polyoma small and middle T antigens and SV40 small t antigen form stable complexes with protein phosphatase 2A," <i>Cell</i> , 60:167-176 (1990)	
	CN1	Perez-Stable, C., <i>et al.</i> , "Prostate Cancer Progression, Metastasis, and Gene Expression in Transgenic Mice," <i>Cancer Res</i> , 57:900-6 (1997).	
	CO1	Rao, K.V., "Structure of Sangivamycin," <i>J Med Chem</i> , 11:939-41 (1968).	
	CP1	Rich, J.N., <i>et al.</i> , "A Genetically Tractable Model of Human Glioma Formation," <i>Cancer Res</i> , 61:3556-60 (2001).	
	CQ1	Richard, D., <i>et al.</i> , "Free radical production and labile iron pool decrease triggered by subtoxic concentration of aclarubicin in human leukemia cell lines," <i>Leukemia Res</i> , 26:927-931 (2002).	
	CR1	Ruediger, R., <i>et al.</i> , "Alterations in protein phosphatase 2A subunit interaction in human carcinomas of the lung and colon with mutations in the A β subunit gene," <i>Oncogene</i> , 20:1892-1899 (2001).	
	CS1	Ruediger, R., <i>et al.</i> , "Disruption of protein phosphatase 2A subunit interaction in human cancers with mutations in the A α subunit gene," <i>Oncogene</i> , 20:10-15 (2001).	
	CT1	Sabatini, D.M., <i>et al.</i> , "RAFT1: A mammalian protein that binds to FKBP12 in a rapamycin-dependent fashion and is homologous to yeast TORs," <i>Cell</i> , 78:35-43 (1994)	
	CU1	Sandmoller, A., <i>et al.</i> , "A Transgenic Mouse Model for Lung Adenocarcinoma," <i>Cell Growth & Differ</i> , 6:97-103 (1995).	
	CV1	Schreiber, S.L., "Chemical Genetics Resulting from a Passion for Synthetic Organic Chemistry," <i>Bioorg. Med. Chem.</i> , 6:1127-1152 (1998).	
	CW1	Sellers, W.R. and Kaelin, W.G., "Role of the retinoblastoma protein in the pathogenesis of human cancer," <i>J Clin Oncol</i> , 15:3301-3312 (1997).	
	CX1	Shawver, L.K., <i>et al.</i> , "Smart drugs: Tyrosine kinase inhibitors in cancer therapy," <i>Cancer Cell</i> , 1:117-123 (2002).	
	CY1	Sherr, C.J., "The <i>INK4a</i> /ARF Network in Tumour Suppression," <i>Nat Rev Mol Cell Biol</i> , 2:731-737 (2001).	
	CZ1	Shi, Y., <i>et al.</i> , "Enhanced Sensitivity of Multiple Myeloma Cells Containing <i>PTEN</i> Mutations to CCI-779," <i>Cancer Res</i> , 62:5027-34 (2002).	
	CA2	Simons, A., <i>et al.</i> , "Establishment of a Chemical Synthetic Lethality Screen in Cultured Human Cells," <i>Genome Res</i> , 11:266-273 (2001).	
	CB2	Stockwell, B. R., "Chemical Genetic Screening Approaches to Neurobiology," <i>Neuron</i> , 36:559-562 (2002).	
	CC2	Stockwell, B. R., "Frontiers in chemical genetics," <i>Trends Biotechnol</i> 18, 449-55, (2000)	
	CD2	Stockwell, B.R., "Chemical Genetics: Ligand-Based Discovery of Gene Function," <i>Nat Rev Genet</i> , 1:116-125 (2000).	
	CE2	Stockwell, B.R., "The biological magic behind the bullets," <i>Nature Biotechnology</i> , 22(1):37-38 (2004).	
	CF2	Stockwell, B.R., <i>et al.</i> , "High-throughput screening of small molecules in miniaturized mammalian cell-based assays involving post-translational modifications," <i>Chem Biol</i> , 6:71-83 (1999).	
	CG2	Testa, J.R. and Giordano, A., "SV40 and cell cycle perturbations in malignant mesothelioma," <i>Seminars in Cancer Biol</i> , 11:31-8 (2001).	
	CH2	Torrance, C.J., <i>et al.</i> , "Use of isogenic human cancer cells for high-throughput screening and drug discovery," <i>Nat Biotechnol</i> , 19:940-945 (2001).	
	CI2	Traganos, F., <i>et al.</i> , "Induction of Apoptosis by Camptothecin and Topotecan," <i>Ann N Y Acad Sci</i> , 803:101-10 (1996).	
	CJ2	Tsao, Y-P., <i>et al.</i> , "Interaction between Replication Forks and Topoisomerase I-DNA	

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